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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
09/282,239	03/31/1999	STEVEN A. GOLDMAN	29556.1540 (SU-1976)	8339
11951 LeClairRyan	7590 03/13/2012 EXAMINER		IINER	
70 Linden Oaks			HUTSON, RICHARD G	
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Please find below and/or attached an Office communication concerning this application or proceeding.

The time period for reply, if any, is set in the attached communication.

1	RECORD OF ORAL HEARING
2	INVESTIGATION OF A TEXAS AND TRANSPORT OF THE
3	UNITED STATES PATENT AND TRADEMARK OFFICE
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5	DEFORE THE DOADD OF DATENT ADDEALG
6 7	BEFORE THE BOARD OF PATENT APPEALS
8	AND INTERFERENCES
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10	Ex Parte STEVEN A. GOLDMAN and SU WANG
11	Ex Parte STEVEN A. GOLDMAN and SU WANG
12	
13	A1 2011 012422
13	Appeal 2011-012422
15	Application 09/282,239 Technology Center 1600
16	Technology Center 1600
17	
18	Oral Hearing Held: January 19, 2012
19	Ofai freating field. January 19, 2012
20	
21	Before TONI R. SCHEINER, ERIC B. GRIMES, and
22	STEPHEN WALSH, Administrative Patent Judges.
23	51E111E1 WALSII, Administrative I dieni Judges.
24	APPEARANCES:
25	THILDING HOLD.
26	ON BEHALF OF THE APPELLANT:
27	ON BEHALF OF THE ALTEREMAN.
28	MICHAEL L. GOLDMAN, ESQUIRE
29	LeClairRyan
30	70 Linden Oaks
31	Suite 210
32	Rochester, New York 14625
33	10000000, 1.0.0. 1.0.0.
34	The above-entitled matter came on for hearing on Thursday,
35	January 19, 2012, at the U.S. Patent and Trademark Office, 600 Dulany
36	Street, Alexandria, Virginia, before David Voigtsberger, a Notary Public
37	

1	<u>PROCEEDINGS</u>
2	THE USHER: Calendar No. 79, Appeal No. 2011-012422,
3	Mr. Goldman.
4	JUDGE SCHEINER: Thank you. Good afternoon. Mr. Goldman,
5	are you the Inventor?
6	MR. GOLDMAN: I am not.
7	JUDGE SCHEINER: Oh, okay.
8	MR. GOLDMAN: I am Michael Goldman, and the Inventor is Steven
9	Goldman.
0	JUDGESCHEINER: I see. Oh, yes. Now I see that. Okay. Before
1	you get started, I wanted to let you know that Judge Walsh is joining us from
2	a remote site, and
3	MR. GOLDMAN: Good afternoon.
4	JUDGE WALSH: Good afternoon.
5	COURT REPORTER: And if you have a business card, I'd
6	appreciate it.
7	MR. GOLDMAN: Sure. Thank you.
8	JUDGE SCHEINER: And before you do get started, I would like to
9	suggest that you spend the bulk of your time on the art rejection.
20	MR. GOLDMAN: Certainly.
21	JUDGE SCHEINER: Okay.
22	MR. GOLDMAN: Okay. Well, good afternoon. I'm here to discuss
23	the appeal from the Examiner's final rejection. Subject matter is an enriched
24	or purified preparation of human mitotic oligodendrocyte progenitor cells.

This is a collection of cells that are in the differentiation scheme between 1 2 stem cells and between fully differentiated oligodendrocytes. 3 The important thing about the claims is -- every limitation is 4 important, but the key ones. I would say, are the fact that these are human 5 cells and that they differentiate preferentially to oligodendrocytes. 6 JUDGE SCHEINER: Okay. Can we -- can I interrupt you right 7 there? 8 MR. GOLDMAN: Certainly. 9 JUDGE SCHEINER: I just want to make sure that we're all 10 understanding the claims the same way. This is a population of cells that 11 will -- the majority of which will mature into oligodendrocytes, if and when 12 they're cultured in the manner discussed in the body of the claim. 13 MR GOLDMAN: That's correct 14 JUDGE SCHEINER: Okay. But it's the actual, the population that's 15 being claimed, not the treated -- or, well, for lack of a better word, the 16 subsequently cultured population. 17 MR. GOLDMAN: It's the oligodendrocyte progenitor cell 18 population, not the oligodendrocytes that we're claiming. 19 JUDGE SCHEINER: No, no, no. I understand that, but I just want to 20 understand that presumably, if this population of cells is cultured in 21 something else, you don't necessarily get the majority maturing into 22 oligodendrocytes. Is that a fair --23 MR. GOLDMAN: I'm not sure about that. 24 JUDGE SCHEINER: Okav.

1 MR. GOLDMAN: It could be that they differentiate in some other 2 way, and get some other result, is the idea. 3 JUDGE SCHEINER: Okav. MR. GOLDMAN: So we wanted to be clear that it was using that 4 5 culture condition. 6 JUDGE SCHEINER: Okay. Because when we start talking about 7 example 7in the reference, I'm going to have a couple questions about that. 8 MR. GOLDMAN: I imagine you would. 9 JUDGE GRIMES: While we're still looking at the claim, is there a 10 difference between 04-positive oligodendrocytes and galactocerebroside-11 positive oligodendrocytes and oligodendrocytes? 12 MR. GOLDMAN: I think it depends on what the -- I mean, the 13 answer is ves. I think it depends on the source, and this is probably pretty key to the idea of rats versus humans. So for example, you know, this has 14 15 been our argument, is rats are different than -- here you go. Besides their 16 being different, the cells are different as well. So in rats, the 17 oligodendrocytes are mitotic. They produce, they divide, and the oligos and 18 the -- the oligodendrocytes themselves and the progenitors will express as a 19 04 marker. But on the other hand, when you look at humans, only the 20 oligodendrocyte progenitors -- or, the oligodendrocyte -- yeah, only the 21 oligodendrocytes express that, but not the progenitors. JUDGE SCHEINER: Can I interrupt you again? 22 23 MR. GOLDMAN: Sure. JUDGE SCHEINER: Rao -- am I pronouncing that correctly? Rao? 24 25 MR. GOLDMAN: Rao. Yes.

1 JUDGE SCHEINER: In figure 1, their cell that they've labeled 14. 2 which is progenitor of oligodendrocytes and astrocytes is labeled 04-3 negative. Now, I understand this is just a diagram, but --4 MR. GOLDMAN: And it's also modeled on rats. 5 JUDGE SCHEINER: Right, I understand that, But I -- didn't I just 6 understand you to say that rat progenitors are 04-positive? 7 MR. GOLDMAN: Let me see. Yeah, the problem is 14 is not rat progenitor. It's a less-differentiated state of cells. So as you can see from 8 9 the diagram in figure 1, those cells can either differentiate to astrocytes or 10 oligodendrocytes. And in fact, this is --11 JUDGE SCHEINER: Okav. I know we're jumping around a lot --12 MR. GOLDMAN: That's okay. 13 JUDGE SCHEINER: But, you know, your claim seemed to be based 14 on the results that you got in example 6 in the specification? Is that correct? 15 MR. GOLDMAN: Certainly --16 JUDGE SCHEINER: I mean, that's one basis we know. 17 MR. GOLDMAN: Yeah. Certainly. 18 JUDGE SCHEINER: Okay. And example 6, there were some 19 astrocytes, and even some neurons observed. 20 MR. GOLDMAN: That's right. But we're not saving there are no 21 astrocytes. That's the majority limitation. 22 JUDGE SCHEINER: Okav. Right. 23 MR. GOLDMAN: So the difference between our claims and what 24 Rao is doing is that the majority are actually astrocytes.

JUDGE SCHEINER: Okay. That's one thing. Can you show us 1 2 where there's evidence of that in -- that Rao got a majority of astrocytes? 3 MR. GOLDMAN: Sure. Examples 14 and 15 of *Rao* in the 4 reference. 5 JUDGE SCHEINER: Okay. Let me get there. Sorry, I know we're 6 clicking all over the place up here. We have all this on the screens up here. 7 MR. GOLDMAN: I do it the old-fashioned way. 8 JUDGE SCHEINER: You said 14 and 15? 9 MR. GOLDMAN: Yeah. 10 JUDGE SCHEINER: Okay. Well, let me make sure I understand the 11 earlier examples first. And I know I'm interrupting your flow here, but I do 12 have questions. 13 MR. GOLDMAN: I'm here to explain the case to you, for your 14 auestions. 15 JUDGE SCHEINER: Okay, the first several examples in Rao, it 16 looks like they're all the same cells, and what they did was they cultured 17 them after the initial isolation from -- was it neural tube? 18 MR. GOLDMAN: Of fetal rats. That's right. 19 JUDGE SCHEINER: Fetal rats, yes. I do know that they were rats. 20 The next several examples are taking those same cells and culturing them 21 under different conditions and under 1, if I understand the examples 22 correctly, under example 2, they culture them with certain factors and they 23 get -- well, let's see -- in one case, they get neurons. In one case, they get 24 predominantly astrocytes, and in one case, they get -- yes, in example 5, they 25 get neurons. Example 6, they get -- seems to be astrocytes?

- 1 MR. GOLDMAN: Yeah. I mean, I don't know if they really -- they 2 talk about all of this, but I don't think they get into, you know, quantifying 3 the yield of one or the other. 4 JUDGE SCHEINER: Exactly. And that's why -- that's where my 5 question comes in, because in example 7, they get -- they, under certain 6 culture conditions, they seem to get predominantly -- I mean, or at least --7 it's true that they don't -- there are no percentages here, but they're focusing 8 on the oligodendrocytes. 9 MR. GOLDMAN: But see, the thing is --JUDGE SCHEINER: So how are examples 14 and 15 different? 10 11 MR. GOLDMAN: Well, the first thing I'd like to point out is that 12 Dr. Rao has put in two declarations in support of our patentability 13 JUDGE SCHEINER: No. I understand that. 14 MR, GOLDMAN: And Dr. Rao, in that, has pointed out that he 15 regards his method as astrocyte-biased, as far as he's -- progenitor cells, 14. 16 JUDGE SCHEINER: Right. But I was -- we were looking for 17 evidence in those declarations. 18 MR. GOLDMAN: Right. Well, I think I would read 7 as just silent on
- 20 JUDGE SCHEINER: Okay.
- 21 MR. GOLDMAN: I don't think 7 speaks to it one way or the other,
- 22 whereas 14 and 15 very clearly do.
- JUDGE SCHEINER: Okay. So let's -- if you could walk us through
- 24 14 and 15, then.

the issue.

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1 MR. GOLDMAN: Sure. So, well, I'll just go -- on 14, if you go just 2 to the conclusion, I mean, it says that the -- it's the last, in column 20. 3 paragraph -- well, yeah, the paragraph that begins on line 53 and it concludes 4 with the culture consists of 30 percent astrocyte and 50 percent -- or, I'm sorry, 30 percent oligodendrocytes, 50 percent astrocytes, and 20 percent 5 other, so -- and they're talking about these A2B5 -- I believe what they're 6 7 referring to -- it's not, you know, as clear as it could be, but I believe what 8 they're talking about is that in the context of the drawing of figure 1, nothing 9 else makes sense -- that what happens to 14 when they culture under these 10 conditions is they get a preference for astrocytes, which is not what we want 11 and not what we claim. 12 JUDGE SCHEINER: And then the next? 13 MR. GOLDMAN: And example 15 is, this is actually if you look in 14 column 22, lines 9 through the end of that paragraph, it says the result 15 suggested that in the presence of these agents, the A2B5 cells predominantly 16 differentiated the cells with a type-2 astrocyte, astrocyte phenotype. 17 JUDGE SCHEINER: The CNTF bFGF, are those -- then you've got 18 FGF-2 in the claim. They're not -- you know, they're not --19 MR. GOLDMAN: I mean, you know, they are identical cultural 20 conditions, but on the other hand, there's nothing to indicate that something 21 will change here from what they're teaching based on the culture conditions. 22 And again, Dr. Rao has testified that these cells are astrocytic bias, they have 23 an astrocytic bias. And that's not what we want. 24 JUDGE SCHEINER: These are the same -- so 14 and 15 start out 25 with the same cells that examples 5, 6, and 7 do?

MR. GOLDMAN: That's my read on it. 1 2 JUDGE SCHEINER: Okav. But don't examples 5, 6, and 7 show 3 that the culture conditions are critical? That you do get different --MR. GOLDMAN: Well, I guess what I view them as is they're just 4 5 studying the different aspects of what's shown in figure one, as far as these different parts of the differentiation scheme. So I don't know that they're 6 7 doing anything other than going through and showing what happened at each 8 one of these different stages, and then finally, when they get around to 14 9 and 15, they studied, you know, the actual bias of these bipotential cells. 10 JUDGE SCHEINER: Okay. Can we go back for a minute now and 11 just look at now -- okay, our claim is human cells. 12 MR. GOLDMAN: Right. 13 JUDGE SCHEINER: Rao is rat cells. 14 MR. GOLDMAN: True. 15 JUDGE SCHEINER: And there is a mention in there that this 16 procedure of theirs, where they're taking -- they're teasing cells out of neural 17 crest and then doing, you know, clonal so on and so forth. 18 MR. GOLDMAN: Right. 19 JUDGE SCHEINER: They're not doing a FACS, a fluorescence-20 activated cell sorting procedure. But there is a suggestion in here that you 21 could take Rao's -- that Rao suggests that you could take this procedure and 22 apply it to human cells. 23 MR. GOLDMAN: Okav. I guess I would differ with that. I think, 24 vou know, the Examiner has certainly seized on, and I guess it's column 6 --

1 JUDGE SCHEINER: So I guess what we'd like to focus on is -- I 2 mean, I understand that your -- correct me if I'm wrong, but I think your 3 position, in part, is that you don't have an actual anticipation because of --4 because this is rat, not human, and that the reference is not enabling for 5 human. 6 MR. GOLDMAN: That's right. 7 JUDGE SCHEINER: Can you focus on the evidence that establishes 8 that -- that you feel establishes that this is not an enabling reference? 9 MR. GOLDMAN: Not enabling for human? 10 JUDGE SCHEINER: For human, ves. 11 MR. GOLDMAN: Yeah. I mean, I don't have any issue with the ref. 12 JUDGE SCHEINER: Right. 13 MR. GOLDMAN: I mean, the only -- basically, all the work is done 14 with rats. There's a slight mention of use in human, but really, when you 15 read it, it's just about these stem cells that are upstream of the cells that have 16 the bipotential differentiation. So it's not really clear what they're trying to 17 say, but I think the evidence that we've put into the record is that rats and 18 humans are pretty different. And I believe the declarations of Dr. Goldman speak to that. 19 20 JUDGE SCHEINER: Okay. We have at least two, is that correct? 21 MR. GOLDMAN: There's two. It's the first, and the third. And I 22 believe they're both in paragraph 7. 23 JUDGE SCHEINER: Okav. 24 MR. GOLDMAN: Of each. So the -- in those declarations -- and I 25 could pull them out and read them if you want -- but basically, they talk

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- 1 about how the rat oligos themselves, oligodendrocytes, are mitotic and the 2 human oligodendrocytes aren't and that the rat oligodendrocytes and their 3 progenitors express the same markers, while the human ones don't. 4 JUDGE SCHEINER: But that's actually not consistent with Rao, is 5 it? Well, at least the earlier cell that they had. 6 MR. GOLDMAN: Well, see, but I don't think what we're talking 7 about there is what Rao really deals with. And what Rao -- that, yeah, I 8 don't think --9 JUDGE SCHEINER: Oh, no, I understand that's your position. I'm 10 just looking for the evidence --11 MR. GOLDMAN: 14 is not an oligodendrocyte progenitor, except in
- 11 MR. GOLDMAN: 14 is not an oligodendrocyte progenitor, except in
 12 the broadest -- in the way we're claiming it. It's a progenitor that will
 13 actually differentiate either to astrocytes or oligos, but it -- with the bias
 14 being to the astrocytes. And so what we're talking about is something that -15 and it's claimed, I think, that our progenitors --
 - JUDGE SCHEINER: Oh, no, I do understand that. I'm just looking for some hard evidence.
 - MR. GOLDMAN: Okay. Okay, well I think the declaration does point out that there are profound differences between humans and rats, and the idea that one would be able to go from results with humans, given the ---from rats, given the differences that are presented in the declaration to rats is pretty tough to fathom.
- I mean, the evidence also is that this was quite a big event when our
 Inventors did this. It's published in *Nature Medicine* as being the first time
 that the human oligodendrocyte progenitor cells were isolated, and there was

- 1 much talk about why that was scientifically interesting for that reason, but 2 also therapeutically interesting in that it would provide a vehicle to treat 3 such diseases as multiple sclerosis and other diseases involving 4 demyelination or the inability to produce it. 5 So I think the evidence -- there's really not much evidence to support the Examiner's position, frankly, on this, other than he tries to kind of tie 6 7 well, this little mention of human over here, and then it's generic in the 8 drawings, and then it got this rat cell stuff that should just be put together. 9 And I just don't think there's any evidence to support the position that
- 10 humans -- or rat work is germane to what you'd expect with human work. 11 And I have a problem -- you know, the biggest problem I think we've had is 12 what kind of reference is this reference? Is it an anticipatory reference, or is 13 it one of obviousness? I think --
- 14 JUDGE SCHEINER: Is it -- oh, it's actually applied 102, 103. Is that 15 correct?
- 16 MR. GOLDMAN: Well, he ties -- yeah. And the problem is that 17 okay, maybe when you look at the general, you know, at the mention of 18 humans, you could say okay, well, that's about humans. But everything else 19 is either generic or it's about rats. And I think once you move to the 20 examples, we're talking -- I don't see how that reference continues to be an 21 anticipatory reference, because the data is all about rats, and the evidence is 22 that rats are, you know, again -- it sounds silly -- are different than humans.
- 23 JUDGE SCHEINER: Right.

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MR. GOLDMAN: Sorry. So, I, you know, I think we put a lot of evidence into the record based on what we were faced with at the -- during

- 1 prosecution to demonstrate that this was not an appropriate rejection. And
- 2 again, Dr. Rao also supports the idea that this is a very different
- 3 phenomenon in that we're getting the bias to the oligodendrocytes, which is
- 4 what you need to be able to treat these diseases of demyelination. Having an
- 5 astrocytic bias doesn't help you, in that regard.
- $\label{eq:JUDGE SCHEINER: I'm sorry that I sort of jumped in, and I'm sure} \\$
- 7 you had some kind of a presentation. Would you like to give it now, or do
- 8 you --
- 9 MR. GOLDMAN: I'll just say a few more things.
- 10 JUDGE SCHEINER: Okay.
- 11 MR. GOLDMAN: But basically, you've, you know, it was fine.
- 12 JUDGE SCHEINER: Okay.
- 13 MR. GOLDMAN: One word about -- one word, or a few words about 14 the 112 rejection.
- 15 JUDGE SCHEINER: Yes, Okav.
- MR. GOLDMAN: Basically, I think we've said our piece on it for the
- most part, but it seems like the issue is that we have the word
- 18 oligodendrocyte in there. Well, the record is pretty clear on -- and I can
- 19 point you to evidence of that -- that these 04 cells in the context that they
- 20 were doing that experimental work, were oligodendrocytes. And again, the
- 21 Goldman declaration, paragraph seven, the Goldman, the third one, said,
- 22 points out that these --
- JUDGE SCHEINER: Third? Oh, I'm sorry. I seem to only have two
- 24 Goldman declarations.
- 25 MR. GOLDMAN: Oh, there was three.

3 JUDGE SCHEINER: Oh, it's called the third. Okav. Oh, yeah. I'm 4 sorry. We do have --5 MR. GOLDMAN: Because I don't know if the second one actually 6 is --7 JUDGE SCHEINER: We do have that. Sorry. 8 MR. GOLDMAN: Yeah. So I think it was, like, in paragraph seven 9 when we talk about how the oligodendrocytes are recognized by antibodies 10 that have this, you know, are looking for this 04 marker. So I think the idea 11 that --12 JUDGE SCHEINER: Well, I think the Examiner in the response 13 mentioned that Schwann cells also have 04 --14 MR. GOLDMAN: But in the context of what we're disclosing, it's 15 not just the general -- so I think in the context of the basis for that argument. 16 I think we're in pretty good shape as far as not being new matter. The other 17 thing I wanted to point -- a couple things I wanted to point out. If, first of all, if you agree with us on those claims with the percentages of the -- well, 18 19 claims 42 to 44 --20 JUDGE SCHEINER: You know we don't tell you anything around 21 here. 22 MR. GOLDMAN: I know. But if you, you know, agree that there's 23 not a 112 problem, I don't see how *Rao* gets close to that. They're talking 24 about an astrocytic bias, and we're talking about a higher percentage of 25 claims being -- a specific percentage being oligodendrocytes. And the only

JUDGE SCHEINER: I will go into the record and --

MR. GOLDMAN: One's called the third. One's called the third.

1 other thing I wanted to mention was that we think the human fetal distinction 2 is also worth paying attention to. 3 JUDGE SCHEINER: Fetal versus adult, you mean? 4 MR. GOLDMAN: Yes. Sorry. Thank you. So basically, we have 5 certain claims that are limited to adult, and we think we presented evidence 6 that adults are different -- again, silly sounding -- but adults are different 7 than fetal, so the extent you're looking at Rao, it's fetal, and we don't see 8 how that is a basis to reject the adult, even if you think the examples would 9 otherwise be suitable to reject our more generic claims. And I think we decided to -- Dr. Rao's first declaration, paragraph 10 11 eight, where we pointed out that there are, you know, significant differences 12 between the two. And again, if that's something that you agree with us on, 13 claims 26 and 42, 43, would be patentable regardless of how you rule on the 14 others. So I don't have --15 JUDGE SCHEINER: Okav. 16 MR. GOLDMAN: I got my piece in from your -- from answering 17 your questions, so --18 JUDGE SCHEINER: All right. Well, I don't pretend to understand 19 this case completely yet, but I will before we --20 MR. GOLDMAN: Okay. Well, thank you for your time. 21 JUDGE SCHEINER: Just, I'm sorry. Judge Walsh, did you have any 22 questions? 23 JUDGE WALSH: No. Thank you. 24 MR. GOLDMAN: Okav. Thank you.

JUDGE SCHEINER: Did vou have anything?

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- 1 JUDGE GRIMES: No.
- 2 MR. GOLDMAN: Okay. Thank you very much.
- 3 (Whereupon, the proceedings, at 2:30 p.m., were concluded.)